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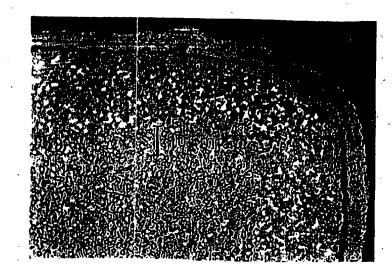
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(54) Title: METHOD OF MAKING CEMENTED CARBIDE WITH BINDER PHASE ENRICHED SURFACE ZONE

(57) Abstract

The present invention relates to method of making a cemented carbide insert, comprising a cemented carbide substrate and a coating. The substrate contains WC and cubic carbonitride phase in a binder phase based of Co and/or Ni and has a binder phase enriched surface zone essentially free of cubic phase. The binder phase enriched surface zone prevails over the edge. By sintering in an atmosphere essentially consisting of nitrogen the thickness of the binder phase enriched zone can be controlled.



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Method of making cemented carbide with binder phase enriched surface zone

The present invention relates to coated cemented carbide inserts with unique edge security in sticky work piece materials such as stainless steel, achieved with a binder phase enriched surface zone extending over the edge.

Coated cemented carbide inserts with binder phase enriched surface zone are today used to a great extent for machining of steel and stainless materials. Thanks to the binder phase enriched surface zone, an extension of the application area for the cutting material has been obtained.

Methods or processes to make cemented carbide containing WC, cubic phase (gamma-phase) and binder phase with binder phase enriched surface zones are known through a number of patents and patent application.

According to e.g. US Patents 4,277,283 and 4,610,931 nitrogen containing additions are used and sintering takes place in vacuum whereas according to US Patent 4,548,786 the nitrogen is added in gas phase. Hereby in both cases a binder phase enriched surface zone essentially free of cubic phase is obtained. US Patent 4,830,934 describes a binder phase enrichment obtained through describes a binder phase enrichment obtained through describes a binder phase enrichment obtained which also contains cubic phase.

It is well known in the art that the thickness of the binder phase enriched zone decreases towards sharp corners, such as the cutting edge of a cutting insert, and that a brittle binder phase depleted zone, enriched in cubic phase, is present in the edge area and often limits the use of binder phase enriched cemented

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carbides especially in work piece materials with high demands on edge toughness.

However, the edges of a cutting insert has to be edge rounded to a certain radius of the order of 50-100 µm or less in order to be useful. The edge rounding is generally made after sintering by an edge rounding operation. In this operation the thin outermost binder phase enriched zone is completely removed and the hard, brittle area is exposed. As a result a hard but brittle edge is obtained resulting in an increased risk for problems with brittleness in the edge particularly in applications demanding high edge toughness.

One method of reducing this drawback of binder phase enriched sintered cemented carbides is described in US 5,484,468. This method is, however, not sufficient in very difficult work piece materials such as austenitic stainless steel and may result in an unwanted decrease in the deformation resistance.

A method of maintaining the binder phase enriched zone in the edge portion of a cemented carbide insert is disclosed in EP-A-0569696. According to this application this effect is obtained if Zr and/or Hf is present in the cemented carbide.

According to Swedish Patent application SE 9501383-5 the thickness of the binder phase enriched surface zone can be maintained over the edge also in cemented carbide free of Hf and Zr if certain conditions are fulfilled particularly with regard to the titanium and nitrogen content within the cubic phase as well as the overall carbon content. A favourable influence on the edge toughness in sticky materials such as austenitic stainless steel can thereby be obtained. However, the binder phase enriched zones according to this application often becomes to deep and difficult to control.

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It is an therefore an object of the present invention to provide a method of making a cemented dearbide insert allowing a better control of the thickness of the binder phase enriched zone.

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According to the presently claimed invention there is provided a cemented carbide insert, comprising a cemented carbide substrate and a coating. The substrate contains WC and cubic carbonitride phase in a binder phase based of Co and/or Ni and has a binder phase enriched surface zone essentially free of cubic phase. The binder phase enriched surface zone prevails over the edge. As a result an insert according to the invention has improved edge toughness and is particularly useful for machining of sticky work piece materials such as stainless steel. (Although the cubic phase is essentially a carbonitride phase the material is herein referred to as a cemented carbide.)

Fig. 1 shows in 800% the binder phase enriched zone under a cutting edge rounded to a 50 μm radius in a coated cemented carbide according to the invention.

It has now surprisingly been found that by performing part of the sintering under nitrogen pressure the thickness of the binder phase enriched surface zone can be controlled with gradient prevailing in the vicinity of the edge.

The invention, thus, relates to a method of making cutting inserts comprising a cemented carbide substrate consisting of a binder phase of Co and/or Ni, WC and a cubic carbonitride phase with a binder phase enriched surface zone essentially free of cubic phase and a coating. A powder mixture containing WC, 6-14 atom-%, preferably 8-11 atom-% binder phase and 3-8 atom-%, preferably 4-6 atom-% of Ti and at least one of Ta and Nb such that the Ti/(Ta+Nb) atomic ratio is >2, preferably >3. Ta and/or Nb is/are added as carbides

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whereas Ti is added as TiC, TiCN and/or TiN in such proportions that the nitrogen content of the carbonitride phase expressed as x in the formula, $(Ti, Nb, Ta)(N_X, C_{1-X})$ shall be >0.2, preferably 0.3-0.4. The powder mixture is mixed with pressing agent and 5 possibly carbon such that the carbon content is 0-0.15, preferably 0.05-0.15, weight-%, above the stoichiometric content and the mixture is milled and dried to obtain a powder material. Next, the powder material is compacted 10 and sintered. During heating to sintering nitrogen gas may be supplied to the furnace at 0-500 mbar, preferably 10-40 mbar, in order to prevent denitrification prior to pore closure at temperatures above 1200°C. Sintering is performed at a temperature of 1380-1520°C, in a protective atmosphere consisting essentially of 15 nitrogen, the nitrogen pressure adjusted to impede gradient growth, with a period of sintering in an atmosphere without nitrogen, the time of this period adjusted to obtain the gradient zone depth desired. The nitrogen pressure required to impede gradient growth 20 depends on composition of the carbide body, sintering temperature and on the furnace used. The time required in atmosphere without nitrogen addition depends on sintering temperature and furnace used. It is within the purview of the skilled artisan to determine whether the 25 requisite binder phase enrichment has been obtained and to modify the sintering conditions in accordance with the present specification, if desired, to effect the desired binder phase enrichment.

Cooling can be performed according to standard practice or as disclosed in US 5,484,468. After conventional post sintering treatments including edgerounding a hard, wear resistant coating according to above is applied by CVD-, PVD- or MT-CVD-technique.

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The present invention also relates to a cutting insert comprising a cemented carbide substrate with a binder phase enriched surface zone and a coating, said substrate consisting of a binder phase of Co and/or Ni, WC and a cubic carbonitride of W , Ti and at least one of the metals Ta, Nb, Mo, V, or Cr with a binder phase enriched surface zone being essentially free of cubic phase.

Preferably the cemented carbide contains 6-14 atom-%, most preferably 8-11 atom-%, binder phase, 3-8 atom-%, most preferably 4-6 atom-%, of Ti and at least one of Ta and Nb and rest WC. The average WC grain size shall be between 1.0 and 4 μm , preferably between 1.5 and 3 μm. The Ti/(Ta+Nb) atomic ratio in the carbonitride phase shall be >2, preferably >3, with a nitrogen content expressed as x in the formula, $(Ti, Nb, Ta)(N_x, C_{1-})$ y) shall be >0.2, preferably between 0.3 and 0.4. The depth of the binder phase enriched surface zone close to the edge increases with increased titanium and nitrogen content within the cubic phase and with increased overall carbon content. The maximum nitrogen content that can be used in practice is mainly limited by the increased tendency for A and B type of porosity with increased nitrogen content. However, the maximum nitrogen content can be extended over the above stated limit if the sintering is performed in an inert atmosphere under high pressure. The maximum carbon content that can be used in practice is mainly limited by an increased tendency for carbon precipitation in the binder phase enriched surface zone, reduced coating adhesion and reduced deformation resistance. The carbon content shall correspond to a C-porosity better than C08, preferably C00 just below carbon saturation.

The thickness of the binder phase enriched surface zone shall be

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> 1. below a flat surface 15-45 μm , preferably 25-35 μ m

- 2. close to a sharp edge, before edge rounding, measured perpendicular to the same flat surface as in 1. above, 0.5-1.2 times the gradient zone in 1. above, preferably 0.67-1.2 times the gradient zone in 1. above.
- 3. at the edge after edge rounding 5-30 μ m, preferably 10-25 µm.

The gradient zone depth close to the edge depends on 10 the geometry, a blunt, 90 degrees or more, geometry giving deeper gradient zones.

Inserts according to the invention shall preferably have a coating of TiC, TiCN and/or TiN with a total coating thickness of 3-10 µm, most preferably 4-8 µm, possibly in combination with an Al₂O₃ coating with a thickness of 1-4 µm, most preferably 1.5-3 µm. Other coatings known in the art can also be used such as single or multiple layers of at least one carbide, nitride, carbonitride, oxide or boride of at least one metal of the groups IVb, VB and VIB of the periodic table and/or aluminium oxide by known CVD-, PVD- or MT-CVD-methods.

Example 1 (according to invention) 25

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From a powder mixture comprising 1.69 weight-% TiC, 1.28 weight-% TiN, 1.21 weight-% TaC, 0.76 weight-% NbC, 7.5 weight-% Co, and balance WC with 0.12 weight-% overstoichiometric carbon content, turning inserts CNMG120408 were pressed. The inserts were sintered with H₂ up to 450°C for dewaxing, further in vacuum to 1200°C, and after that with a protective gas of 40 mbar nitrogen up to 1380° C the furnace was then evacuated and filled with nitrogen to 60 mbar and heated to sintering temperature, 1450°C, and held there for 60 minutes, 35

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during these 60 minutes the furnace was evacuated for 15 min and then refilled with nitrogen.

The structure in the surface of the cutting inserts consisted of a 30 μm thick binder phase enriched zone below the flat flank face with a minimum if 25 μm close to the edge.

Example 2 (according to invention)

From a powder mixture comprising 1.69 weight-% TiC,
1.28 weight-% TiN, 1.21 weight-% TaC, 0.76 weight-% NbC,
7.5 weight-% Co, and balance WC with 0.12 weight-%
overstoichiometric carbon content, turning inserts
CNMG120408 were pressed. The inserts were sintered with
H₂ up to 450°C for dewaxing, further in vacuum to
15 1200°C, and after that with a protective gas of 40 mbar
nitrogen up to 1380°C the furnace was then evacuated and
filled with nitrogen to 200 mbar heated to sintering
temperature, 1450°C, and held there for 10 minutes and
then cooled to 1380°C, evacuated and held at 1380°C for
50 minutes and then cooled.

The structure in the surface of the cutting inserts consisted of a 33 μm thick binder phase enriched zone below the flat flank face with a minimum if 23 μm close to the edge.

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Example 3 (prior art).

From a powder mixture comprising 1.69 weight-% TiC, 1.28 weight-% TiN, 1.21 weight-% TaC, 0.76 weight-% NbC, 7.5 weight-% Co, and balance WC with 0.12 weight-% overstoichiometric carbon content, turning inserts CNMG120408 were pressed. The inserts were sintered with H₂ up to 450°C for dewaxing, further in vacuum to 1200°C, and after that with a protective gas of 40 mbar nitrogen up to 1380°C the furnace was then evacuated and filled with argon to 40 mbar heated to sintering

temperature, 1450° C, and held there for 1 hour and then cooled.

The structure in the surface of the cutting inserts consisted of a 46 μm thick binder phase enriched zone below the flat flank face with a minimum if 30 μm close to the edge.

Example 4 (prior art)

From a powder mixture comprising 1.69 weight-% TiC, 1.28 weight-% TiN, 1.21 weight-% TaC, 0.76 weight-% NbC, 7.5 weight-% Co, and balance WC with 0.12 weight-% overstoichiometric carbon content, turning inserts CNMG120408 were pressed. The inserts were sintered with H₂ up to 450°C for dewaxing, further in vacuum to 1380°C the furnace was then filled with argon to 40 mbar heated to sintering temperature, 1410°C, and held there for 1 hour and then cooled.

The structure in the surface of the cutting inserts consisted of a 40 μm thick binder phase enriched zone below the flat flank face with a minimum if 26 μm close to the edge.

Example 5 (comparative)

From a powder mixture comprising 1.69 weight-% TiC,

1.28 weight-% TiN, 1.21 weight-% TaC, 0.76 weight-% NbC,

7.5 weight-% Co, and balance WC with 0.12 weight-%

overstoichiometric carbon content, turning inserts

CNMG120408 were pressed. The inserts were sintered with

H₂ up to 450°C for dewaxing, further in vacuum to

1200°C, and after that with a protective gas of 40 mbar

nitrogen up to 1380°C the furnace was then evacuated and

filled with nitrogen to 100 mbar heated to sintering

temperature, 1450°C, and held there for 1 hour and then

cooled.

The structure in the surface of the cutting inserts consisted of a 26 μm thick binder phase enriched zone below the flat flank face with a minimum if 12 μm close to the edge.

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Examples 1 and 2 show that it is possible to control the depth of the gradient zone without loosing the desired gradient in the vicinity of the edge as in example 5. Examples 3 and 4 show that the gradient zone may grow excessively without nitrogen addition under a part of the sintering.

Claims

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1. Method of making a cutting insert comprising a cemented carbide substrate with a binder phase enriched surface zone and a coating, said substrate consisting of a binder phase of Co and/or Ni, WC and a cubic carbonitride phase, said binder phase enriched surface zone being essentially free of said cubic carbonitride phase and with an essentially constant thickness around the insert characterised in forming a powder mixture containing WC, 6-14 atom-%, preferably 8-11 atom-%, binder phase and 3-8 atom-%, preferably 4-6 atom-%, of Ti and at least one of Ta and Nb such that the Ti/(Ta+Nb) atomic ratio is >2, preferably >3, Ta and/or Nb being added as carbide and Ti as carbide, nitride and/or carbonitride in such proportions that the nitrogen content of the carbonitride phase expressed as x in the formula, $(Ti,Nb,Ta)(N_x,C_{1-x})$, is >0.2, preferably 0.3-0.4

adding to said powder mixture pressing agent and possibly carbon such that the carbon content is 0-0.15 weight-% above the stoichiometric content

milling and drying the mixture to obtain appowder material

compacting and sintering the pewder material whereby between 1200°C and pore closure nitrogen gas may be supplied to the furnace at 0-500 mbar, preferably 10-40 mbar, after which sintering is performed at a temperature of 1380-1520°C, preferably 1410-1450°C, in a protective atmosphere consisting essentially of nitrogen, the nitrogen pressure adjusted to impede gradient growth, with a period of sintering in an atmosphere without nitrogen, the time of this period adjusted to the gradient zone desired, followed by cooling according to standard practice

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applying conventional post sintering treatments including edgerounding and

forming a hard, wear resistant coating of single or multiple layers of at least one carbide, nitride, carbonitride, oxide or boride of at least one metal of the groups IVb, VB and VIB of the periodic table and/or aluminium oxide by known CVD-, PVD- or MT-CVD-technique.

- 2. Cutting insert for machining of sticky work piece materials such as stainless steel comprising a cemented carbide substrate with a binder phase enriched surface zone and a coating, said substrate consisting of a Co binder phase, WC and a cubic carbonitride phase of W and at least one of the metals Ti, Ta, Nb, Mo, V, or Cr said binder phase enriched surface zone being essentially free of said cubic phase characterised in a thickness of said binder phase enriched surface zone of 15-45 μm on a flat surface of said insert and of 5-30 μm in the cutting edge.
- 3. Cutting insert according to claim 2

 20 characterised in said substrate consisting of 6-14 atom-% binder phase, 3-8 atom-%, preferably 4-6 atom-% of Ti and at least one of Ta and Nb such that the Ti/(Ta+Nb) atomic ratio is >2 and that the nitrogen content of the carbonitride phase expressed as x in the formula, (Ti,Nb,Ta)(N_X,C_{1-x}), is >0.2.
 - 4. Cutting insert according to claim 3 characterised in said Ti/(Ta+Nb) atomic ratio >3.
 - 5. Cutting insert according to claims 3 or 4 characterised in 0.3<x<0.4.
- 6 . Cutting insert according to any of claims 2, 3, 4 or 5 characterised in said coating consisting of at least one of TiC, TiCN or TiN with a total coating thickness of 3-10 μm .

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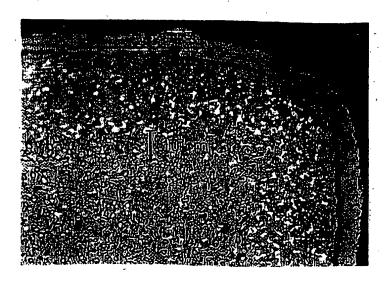


Fig. 1

INTERNATIONAL SEARCH REPORT

International application No.

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CLASSIFICATION OF SUBJECT MATTER IPC6: C22C 29/08 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC6: C22C Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched SE,DK,FI,NO classes as above Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) WPI, PATFULL C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Category* Citation of document, with indication, where appropriate, of the relevant passages X WO 9317140 A1 (SANDVIK AB), 2 Sept 1993 (02.09.93), 1-6 page 2, line 14 - page 6, line 12 X US 4649084 A (THOMAS E. HALE ET AL), 10 March 1987 1-6 (10.03.87), column 2, line 42 - column 3, line 2; column 9, line 19 - column 10, line 56, figure 5 X US 4548786 A (WARREN C. YOHE), 22 October 1985 1-6 (22.10.85), whole document Further documents are listed in the continuation of Box C. See patent family annex. Special categories of cited documents: later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention document defining the general state of the art which is not considered to be of particular relevance "X" document of particular relevance: the claimed invention cannot be erlier document but published on or after the international filing date considered novel or cannot be considered to involve an inventive document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other step when the document is taken alone special reason (as specified) document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination document referring to an oral disclosure, use, exhibition or other being obvious to a person skilled in the art document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of mailing of the international search report Date of the actual completion of the international search 17 -02- 1998 12 February 1998 Authorized officer Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Nils Engnell Facsimile No. +46 8 666 02 86 Telephone No. +46 8 782 25 00

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